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(74) Agent: VERHAGE, Marinus; C.J. van Houtenlaan 36,
NL-1381 CP Weesp (NL).

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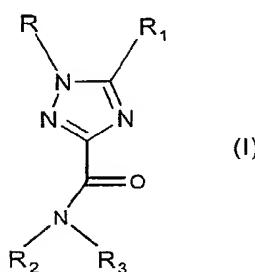
(71) Applicant (*for all designated States except US*): **SOLVAY PHARMACEUTICALS B.V. [NL/NL]**; C.J. van Houtenlaan 36, NL-1381 CP Weesp (NL).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): **LANGE, Josephus H.M. [NL/NL]**; C.J. van Houtenlaan 36, NL-1381 CP Weesp (NL). **KRUSE, Cornelis G. [NL/NL]**; C.J. van Houtenlaan 36, NL-1381 CP Weesp (NL). **McCREARY, Andrew C. [GB/NL]**; C.J. van Houtenlaan 36, NL-1381 CP Weesp (NL). **VAN STUIVENBERG, Herman H. [NL/NL]**; C.J. van Houtenlaan 36, NL-1381 CP Weesp (NL).

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(54) Title: 1H-1,2,4-TRIAZOLE-3-CARBOXAMIDE DERIVATIVES AS CANNABINOID-CB₁ RECEPTOR LIGANDS



(57) Abstract: The present invention relates to a group of 1H-1,2,4-triazole-3-carboxamide derivatives, to methods for the preparation of these compounds, and to pharmaceutical compositions containing at least one of these compounds as an active ingredient. These 1H-1,2,4-triazole-3-carboxamide derivatives are potent cannabinoid-CB₁ receptor agonists, partial agonists, inverse agonists or antagonists, useful for the treatment of disorders involving cannabinoid neurotransmission. The compounds have the general formula (I), wherein R and R₁-R₃ have the meanings given in the specification.

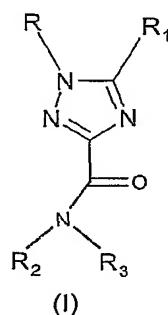
1H-1,2,4- TRIAZOLE-3- CARBOXAMIDE DERIVATIVES AS CANNABINOID-CB₁ RECEPTOR LIGANDS

The present invention relates to a group of 1H-1,2,4-triazole derivatives, to methods
5 for the preparation of these compounds, and to pharmaceutical compositions containing one or more of these compounds as an active ingredient.

These 1H-1,2,4-triazole-carboxamide derivatives are potent cannabinoid-CB₁ receptor agonists, partial agonists, inverse agonists or antagonists, useful for the
10 treatment of psychiatric and neurological disorders, as well as other diseases involving cannabinoid-CB₁ neurotransmission.

1,5-Diaryl-1H-1,2,4-triazole-3-carboxamide derivatives have been described in EP
0346620 and GB 2120665 as herbicides. Recently 1,2,4-triazoles were described as
15 potential agonists and antagonists of cannabinoid-CB₁ and -CB₂ receptors (Jagerovic, N. et al., *Drugs Fut.* 2002, 27(Suppl. A): XVIth Int. Symp. on Medicinal Chemistry, P284)

It has now surprisingly been found that known and new 1,5-diaryl-1H-1,2,4-triazole-3-
20 carboxamide derivatives of the formula (I), as well as prodrugs, salts, and stereoisomers thereof, are potent antagonists, agonists, inverse agonists or partial agonists of the cannabinoid CB₁ receptor:



25

wherein

- R and R₁ independently represent a phenyl, naphtyl, thienyl, pyridyl, pyrimidyl, pyrazinyl, pyridazinyl or triazinyl group, which groups may be substituted with 1-4 substituents X, which can be the same or different, from the group branched or unbranched (C₁₋₃)-alkyl or alkoxy, hydroxy, halogen, trifluoromethyl, trifluoromethylthio, trifluoromethoxy, nitro, amino, mono- or dialkyl (C₁₋₂)-amino, mono- or dialkyl (C₁₋₂)-amido,

30

(C₁₋₃)-alkoxycarbonyl, trifluoromethylsulfonyl, sulfamoyl, (C₁₋₃)-alkylsulfonyl, carboxyl, cyano, carbamoyl, (C₁₋₃)-dialkylaminosulfonyl, (C₁₋₃)-monoalkylamino-sulfonyl and acetyl,

- R₂ represents a hydrogen atom or a branched or unbranched C₁₋₈ alkyl or C₁₈ cycloalkyl-alkyl group or a phenyl, benzyl or phenethyl group which aromatic rings may be substituted with 1-4 substituents X, wherein X has the meaning as indicated above, or R₂ represents a pyridyl or thienyl group,
 - R₃ represents branched or unbranched C₁₋₈ alkyl, C₁₋₈ alkoxy, C₃₋₈ cycloalkyl, C₅₋₁₀ bicycloalkyl, C₆₋₁₀ tricycloalkyl, C₃₋₈ alkenyl, C₅₋₈ cycloalkenyl, which groups may optionally contain one or more heteroatoms from the group (O, N, S), which groups may be substituted with a hydroxy group, an ethynyl group or 1-3 fluoro atoms, or R₃ represents a phenyl, benzyl or phenethyl group which aromatic rings may be substituted with 1-4 substituents X, wherein X has the meaning as indicated above, or R₃ represents a pyridyl, pyrimidyl, pyrazinyl, pyridazinyl, triazinyl or thienyl group which heteroaromatic rings may be substituted with 1-2 substituents X, wherein X has the meaning as indicated above, or R₃ represents a group NR₄R₅ wherein R₄ and R₅, together with the nitrogen atom to which they are bonded, form a saturated or unsaturated, monocyclic or bicyclic, heterocyclic moiety having 4 to 10 ring atoms, which heterocyclic group contains one or two heteroatoms from the group N, O or S, which heteroatoms can be the same or different, which heterocyclic moiety may be substituted with a branched or unbranched C₁₋₃ alkyl, hydroxy or trifluoromethyl group or a fluoro atom, or R₂ and R₃, together with the nitrogen atom to which they are bonded, form a saturated or unsaturated, monocyclic or bicyclic, heterocyclic moiety having 4 to 10 ring atoms, which heterocyclic group contains one or two heteroatoms from the group N, O or S, which heteroatoms can be the same or different, which heterocyclic moiety may be substituted with a branched or unbranched C₁₋₃ alkyl, hydroxy, piperidinyl or trifluoromethyl group or a fluoro atom.
- 30 A group of four 1,5-diaryl-1H-1,2,4-triazole-3-carboxamide derivatives in which the amide N-atom is part of an unsubstituted piperidinyl or morpholinyl group is described by D. Clerin and J.P. Fleury in *Bull. Soc. Chim. Fr.*, 1974, 1-2, Pt.2, 211-217.
- 35 1-(4-Methylphenyl)-5-phenyl-N-(2-pyridyl)-1H-1,2,4-triazole-3-carboxamide is described by M. H. Elnagdi et al. in *Heteroatom Chem.*, 1995, 6, 589-592.
- 40 A group of four 1,5-diaryl-N-(2-pyridyl)-1H-1,2,4-triazole-3-carboxamides is described by A. H. Harhash et al. in *Indian J. Chem.*, 1976, 14B, 268-272.

- Due to the potent cannabinoid-CB₁ receptor agonistic, partial agonistic, inverse agonistic or antagonistic activity the compounds of the invention are suitable for use in the treatment of psychosis, anxiety, depression, attention deficits, memory disorders, cognitive disorders, appetite disorders, obesity, addiction, appetence, drug dependence, neurodegenerative disorders, dementia, dystonia, muscle spasticity, tremor, epilepsy, multiple sclerosis, traumatic brain injury, stroke, Parkinson's disease, Alzheimer's disease, epilepsy, Huntington's disease, Tourette's syndrome, cerebral ischaemia, cerebral apoplexy, craniocerebral trauma, stroke, spinal cord injury, neuroinflammatory disorders, plaque sclerosis, viral encephalitis, demyelinisation related disorders, as well as for the treatment of pain disorders, including neuropathic pain disorders, septic shock, glaucoma, diabetes, cancer, emesis, nausea, gastrointestinal disorders, gastric ulcers, diarrhoea and cardiovascular disorders.
- The affinity of the compounds of the invention for cannabinoid CB₁ receptors was determined using membrane preparations of Chinese hamster ovary (CHO) cells in which the human cannabinoid CB₁ receptor is stably transfected in conjunction with [³H]CP-55,940 as radioligand. After incubation of a freshly prepared cell membrane preparation with the [³H]-ligand, with or without addition of compounds of the invention, separation of bound and free ligand was performed by filtration over glassfiber filters. Radioactivity on the filter was measured by liquid scintillation counting.
- The cannabinoid CB₁ receptor antagonistic, agonistic or partial agonistic activity of compounds of the invention was determined by functional studies using the human CB₁ receptor cloned in Chinese hamster ovary (CHO) cells. CHO cells were grown in a DMEM culture medium, supplemented with 10% heat-inactivated fetal calf serum. Medium was aspirated and replaced by DMEM, without fetal calf serum, but containing [³H]-Arachidonic acid and incubated overnight in a cell culture stove (5% CO₂/95% air; 37 °C; water-saturated atmosphere). During this period [³H]-Arachidonic acid was incorporated in membrane phospholipids. On the test day, medium was aspirated and cells were washed three times using 0.5 ml phosphate-buffered saline, containing 0.2% bovine serum albumin. Stimulation of the CB₁ receptor by WIN 55,212-2 led to activation of PLA₂ followed by release of [³H]-Arachidonic acid into the medium. This WIN 55,212-2-induced release was concentration-dependently antagonized by CB₁ receptor antagonists.
- Cannabinoid agonistic or partial agonistic activity of compounds of the invention can be determined according to published methods, such as assessment of *in vivo* cannabimimetic effects (Wiley, J. L.; Jefferson, R. G; Grier, M. C.; Mahadevan, A.; Razdan, R. K.; Martin, B. R. *J. Pharmacol. Exp. Ther.* 2001, 296 1013).

- The invention relates both to racemates, mixtures of diastereomers and the individual stereoisomers of the compounds having formula (I). Also prodrugs, i.e. compounds which when administered to humans by any known route, are metabolised to 5 compounds having formula (I), belong to the invention. In particular this relates to compounds with primary or secondary amino or hydroxy groups. Such compounds can be reacted with organic acids to yield compounds having formula (I) wherein an additional group is present which is easily removed after administration, for instance, but not limited to amidine, enamine, a Mannich base, a hydroxyl-methylene 10 derivative, an O-(acyloxymethylene carbamate) derivative, carbamate, ester, amide or enaminone. A pro-drug is an inactive compound, which when absorbed is converted into an active form (Medicinal Chemistry: Principles and Practice, 1994, ISBN 0-85186-494-5, Ed.: F. D. King, p. 216).
- 15 The compounds of the invention can be brought into forms suitable for administration by means of usual processes using auxiliary substances and/or liquid or solid carrier materials.

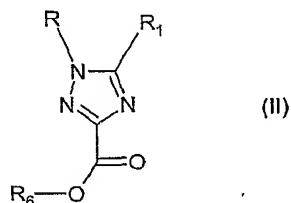
Suitable synthetic routes for the compounds of the invention are the following:

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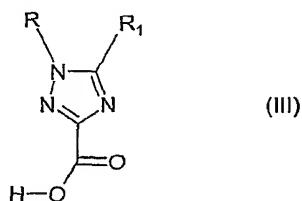
Synthetic route A

Step 1: Ester hydrolysis of a compound having formula (II) wherein R₆ represents a branched or unbranched (C₁₋₄)-alkyl group or a benzyl group,

25



yields a compound having formula (III)



30 wherein R and R₁ have the meanings as described above.

The compounds of the invention having formula (II), wherein R₆ represents a branched or unbranched alkyl group (C₁₋₄) or benzyl group can be obtained according to methods known, for example:

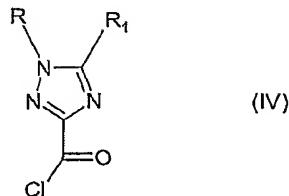
- 5 a) Sawdey, G.W. *J. Am. Chem. Soc.* **1957**, 79, 1955
 b) Czollner, L. *et al.*, *Arch. Pharm. (Weinheim)* **1990**, 323, 225
 c) Eicher, T. and Hauptmann, S. *The Chemistry of Heterocycles*, Thieme Verlag, Stuttgart, 1995 (ISBN 313 100511 4), p. 208-212.
- 10 Step 2: Reaction of a compound having formula (III) with a compound having formula R₂R₃NH wherein R₂ and R₃ have the meanings as described above via activating and coupling methods such as formation of an active ester, or in the presence of a coupling reagent such as DCC, HBTU, BOP, CIP (2-chloro-1,3-dimethylimidazolinium hexafluorophosphate) or PyAOP (7-azabenzotriazol-1-yloxytris(pyrrolidino)phosphonium hexafluorophosphate). Activating and coupling methods of this type are described in
 - a) M. Bodanszky and A. Bodanszky: *The Practice of Peptide Synthesis* Springer-Verlag, New York, 1994; ISBN: 0-387-57505-7;
 - 20 b) K. Akaji *et al.*, *Tetrahedron Lett.* (1994), 35, 3315-3318;
 - c) F. Albericio *et al.*, *Tetrahedron Lett.* (1997), 38, 4853-4856).

This reaction gives a 1H-1,2,4-triazole derivative having formula (I).

25 Synthetic route B

A compound having formula (III) is reacted with a halogenating agent such as thionyl chloride (SOCl₂) or oxalyl chloride. This reaction yields the corresponding carbonyl chloride (acid chloride) (IV).

30



Reaction of a compound having formula (IV) with a compound having formula R₂R₃NH wherein R₂ and R₃ have the meanings as described above gives a

35 1H-1,2,4-triazole derivative having formula (I).

Synthetic route C

- A compound having formula (II) is reacted in an amidation reaction with a compound having formula R_2R_3NH wherein R_2 and R_3 have the meanings as described hereinabove to give a 1H-1,2,4-triazole derivative having formula (I). Such amidation reactions can be promoted by the use of trimethylaluminum $Al(CH_3)_3$ (For more information on aluminum-mediated conversion of esters to amides, see: J. I. Levin, E. Turos, S. M. Weinreb, *Synth Commun.* (1982), 12, 989-993.)

10 **Example I**

Part A: To a stirred solution of dimethyl aminomalonate hydrochloride (25 gram, 0.136 mol) in dichloromethane (200 mL) triethylamine (41.4 mL, 2.2 molar equivalent) is added at 0 °C. 4-Chlorobenzoyl chloride (23.8 gram, 0.136 mol) is slowly added and the resulting solution is allowed to stand at room temperature overnight. Water is added and the organic layer is separated. The water layer is extracted twice with dichloromethane. The collected organic layers are washed with water, dried over $MgSO_4$, filtered and concentrated in vacuo. The residue is recrystallised from methanol (400 mL) to give dimethyl 2-(4-chlorobenzoylamino)malonate (30.5 gram, 79 % yield). Melting point: 146-148°C. 1H -NMR (200 MHz, $CDCl_3$): δ 3.86 (s, 6H), 5.38 (d, J = 6 Hz, 1H), 7.15 (br d, J ~ 6 Hz, 1H), 7.43 (d, J = 8 Hz, 2H), 7.79 (d, J = 8 Hz, 2H).

Part B: To a stirred suspension of 2,4-dichloroaniline (19.44 gram, 0.12 mol) in concentrated HCl (25 mL) and acetic acid (75 mL) at 0 °C is added a solution of $NaNO_2$ (9.0 gram, 0.13 mol) in water (50 mL) and the resulting solution is stirred for 15 minutes. A solution of dimethyl 2-(4-chlorobenzoylamino)-malonate (28.55 gram, 0.10 mol) in acetone (200 mL) is slowly added while keeping the temperature below 0 °C. A solution of K_2CO_3 (120 gram) in water (200 mL) is slowly added and the resulting black mixture is stirred for 30 minutes at 0°C. The mixture is extracted three times with EtOAc. The collected organics are washed with water, aqueous $NaHCO_3$ and water, respectively, dried over $MgSO_4$, filtered and concentrated in vacuo. The residue is dissolved in methanol (500 mL) and a solution of sodium (1 gram) in methanol (75 mL) is added. The resulting stirred mixture is allowed to stand overnight at room temperature and cooled in a refrigerator. The formed precipitate is collected by filtration and washed with methanol to give methyl 5-(4-chlorophenyl)-1-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carboxylate (11.4 gram, 30 % yield). Melting point: 153-154 °C. 1H -NMR (200 MHz, $CDCl_3$): δ 4.07 (s, 3H), 7.28-7.60 (m, 7H).

Part C: To a stirred suspension of methyl 5-(4-chlorophenyl)-1-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carboxylate (11.3 gram, 0.0295 mol) in methanol (100 mL) is added KOH (45 % aqueous solution, 7.5 mL) and the resulting mixture is heated at reflux temperature for 4 hours. The mixture is concentrated in vacuo and water (150 mL) and concentrated HCl are added. The yellow precipitate is collected by filtration, washed with water and dried in vacuo to give 5-(4-chlorophenyl)-1-(2,4-

dichlorophenyl)-1H-1,2,4-triazole-3-carboxylic acid (10.0 gram, 92 % yield). Melting point: 141-144 °C (decomposition).

Part D: To a stirred solution of 5-(4-chlorophenyl)-1-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carboxylic acid (1.48 gram, 4.0 mmol) in acetonitrile (20 mL) is successively added diisopropylethylamine (DIPEA) (1.5 mL, 2.1 molar equivalent), O-benzotriazol-1-yl-N,N,N',N'-tetramethyluronium hexafluorophosphate (HBTU) (1.66 gram, 1.1 molar equivalent) and 1-aminopiperidine (0.44 gram, 1.1 molar equivalent). After stirring overnight an aqueous NaHCO₃ solution is added. The resulting mixture is three times extracted with dichloromethane. The combined organic layers are washed with water, dried over Na₂SO₄, filtered and concentrated *in vacuo* to give a crude oil (3.6 gram). This oil is further purified by flash chromatography (silica gel; EtOAc / petroleum ether (40-60 °C) = 7/3 (v/v)). The purified material is treated with ethanolic HCl (1M solution) to give 5-(4-chlorophenyl)-1-(2,4-dichlorophenyl)-N-(piperidin-1-yl)-1H-1,2,4-triazole-3-carboxamide hydrochloride (1.50 gram, 77 % yield). Melting point: 238-240 °C (decomposition). ¹H-NMR (400 MHz, DMSO-d₆): δ 1.46-1.54 (m, 2H), 1.78-1.85 (m, 4H), 3.22-3.28 (m, 4H), 7.50 (s, 4H), 7.70 (dd, J = 8 and 2 Hz, 1H), 7.85-7.87 (m, 1H), 7.91 (d, J = 8 Hz, 1H), (NH not visible).

Analogously were prepared the examples 2-18:

- 20 2. 5-(4-Chlorophenyl)-1-(2,4-dichlorophenyl)-N-(pyrrolidin-1-yl)-1H-1,2,4-triazole-3-carboxamide hydrochloride. Melting point: 248-255 °C (decomposition).
3. 5-(4-Chlorophenyl)-N-cyclohexyl-1-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carboxamide. Melting point: 186-188 °C.
- 25 4. N-t-Butoxy-5-(4-chlorophenyl)-1-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carboxamide. Melting point: 150-152 °C.
5. 5-(4-Chlorophenyl)-1-(2,4-dichlorophenyl)-N-(n-pentyl)-1H-1,2,4-triazole-3-carboxamide. ¹H-NMR (400 MHz, CDCl₃): δ 0.92 (t, J = 7 Hz, 3H), 1.35-1.44 (m, 4H), 1.62-1.70 (m, 2H), 3.48-3.56 (m, 2H), 7.20-7.25 (m, 1H), 7.34 (dt, J = 8 and 2 Hz, 2H), 7.42-7.50 (m, 4H), 7.54 (d, J = 2 Hz, 1H).
- 30 6. 5-(4-Chlorophenyl)-1-(2,4-dichlorophenyl)-N-(morpholin-4-yl)-1H-1,2,4-triazole-3-carboxamide. Melting point: 184-186 °C.
7. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(piperidin-1-yl)-1H-1,2,4-triazole-3-carboxamide hydrochloride. Melting point: 234-237 °C (decomposition).
- 35 8. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(pyrrolidin-1-yl)-1H-1,2,4-triazole-3-carboxamide hydrochloride. Melting point: 234-236 °C (decomposition).
9. 1-(4-Chlorophenyl)-N-cyclohexyl-5-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carboxamide. ¹H-NMR (400 MHz, CDCl₃): δ 1.14-1.81 (m, 8H), 2.02-2.10 (m, 2H), 4.00-4.11 (m, 1H), 7.08 (br d, J ~ 7 Hz, 1H), 7.26 (br d, J ~ 8 Hz, 2H), 7.34 (br d, J ~ 8 Hz, 2H), 7.40 (dd, J = 8 and 2 Hz, 1H), 7.44-7.48 (m, 2H).

10. N-t-Butoxy-1-(4-chlorophenyl)-5-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carboxamide. $^1\text{H-NMR}$ (400 MHz, CDCl_3): δ 1.38 (s, 9H), 7.25 (br d, $J \sim 8$ Hz, 2H), 7.35 (br d, $J \sim 8$ Hz, 2H), 7.41 (dd, $J = 8$ and 2 Hz, 1H), 7.44-7.48 (m, 2H), 9.18, br s, 1H).
- 5 11. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(n-pentyl)-1H-1,2,4-triazole-3-carboxamide. $^1\text{H-NMR}$ (400 MHz, CDCl_3): δ 0.91 (t, $J = 7$ Hz, 3H), 1.35-1.41 (m, 4H), 1.60-1.70 (m, 2H), 3.48-3.56 (m, 2H), 7.21 (br t, $J \sim 7$ Hz, 1H), 7.26 (br d, $J \sim 8$ Hz, 2H), 7.34 (br d, $J \sim 8$ Hz, 2H), 7.40 (dd, $J = 8$ and 2 Hz, 1H), 7.44-7.48 (m, 2H).
- 10 12. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(morpholin-4-yl)-1H-1,2,4-triazole-3-carboxamide hydrochloride. Melting point: 224-226 °C.
13. 1-(2,4-Dichlorophenyl)-5-(pyridin-2-yl)-N-(piperidin-1-yl)-1H-1,2,4-triazole-3-carboxamide. Melting point: 191-193 °C.
14. 5-(2,4-Dichlorophenyl)-N-(piperidin-1-yl)-1-(4-(trifluoromethyl)phenyl)-1H-15 1,2,4-triazole-3-carboxamide. Melting point: 159-161 °C.
15. 1'-[5-(2,4-dichlorophenyl)-1-(4-(trifluoromethyl)phenyl)-1H-1,2,4-triazole-3-yl]carbonyl]piperidine. Melting point: 155-156 °C.
16. 1-(2,4-Dichlorophenyl)-N-(piperidin-1-yl)-5-(pyridin-3-yl)-1H-1,2,4-triazole-3-carboxamide. Melting point: 219 °C.
- 20 17. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(5,5,5-trifluoropentyl)-1H-1,2,4-triazole-3-carboxamide. $^1\text{H-NMR}$ (400 MHz, CDCl_3): δ 1.63-1.80 (m, 4H), 2.06-2.22 (m, 2H), 3.54 (q, $J \sim 7$ Hz, 2H), 7.26 (m, 3H), 7.34 (br d, $J \sim 8$ Hz, 2H), 7.40 (dd, $J = 8$ and 2 Hz, 1H), 7.44-7.48 (m, 2H).
- 25 18. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(5-fluoropentyl)-1H-1,2,4-triazole-3-carboxamide. $^1\text{H-NMR}$ (400 MHz, CDCl_3): δ 1.63-1.80 (m, 4H), 2.06-2.22 (m, 2H), 3.54 (q, $J \sim 7$ Hz, 2H), 7.22-7.28 (m, 3H), 7.34 (br d, $J \sim 8$ Hz, 2H), 7.40 (dd, $J = 8$ and 2 Hz, 1H), 7.44-7.48 (m, 2H).

Example 19

- 30 Part A: 1-(Chlorophenyl)-5-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carboxylic acid was prepared analogously to the procedure as described in Example 1, Part A-C by using dimethyl aminomalonate hydrochloride, 2,4-dichlorobenzoyl chloride and 4-chloroaniline as starting materials, respectively. Melting point: 102-104 °C. $^1\text{H-NMR}$ (400 MHz, DMSO-d_6): δ 7.36 (br d, $J \sim 8$ Hz, 2H), 7.50 (br d, $J \sim 8$ Hz, 2H), 7.59 (dd, $J = 8$ and 2 Hz, 1H), 7.70 (d, $J = 2$ Hz, 1H), 7.75 (d, $J = 8$ Hz, 1H), OH proton is part of water peak at δ 3.4. Analogously was 1-(chlorophenyl)-5-(2,5-dichlorophenyl)-1H-1,2,4-triazole-3-carboxylic acid prepared by using dimethyl aminomalonate hydrochloride, 2,5-dichlorobenzoyl chloride and 4-chloroaniline as starting materials, respectively.
- 35 Melting point: 183-188 °C. $^1\text{H-NMR}$ (400 MHz, DMSO-d_6): δ 7.41 (br d, $J \sim 8$ Hz, 2H), 7.52 (br d, $J \sim 8$ Hz, 2H), 7.56 (d, $J = 8$ Hz, 1H), 7.65 (dd, $J = 8$ and 2 Hz, 1H), 7.88 (d, $J = 2$ Hz, 1H), OH proton is part of water peak at 83.5.

Part B: To a stirred solution of 1-(chlorophenyl)-5-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carboxylic acid (0.37 g, 1.00 mmol) in dichloromethane (10 mL) is added oxalyl chloride (0.254 g, 2.00 mmol). The resulting mixture is concentrated in vacuo to give crude 1-(chlorophenyl)-5-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carbonyl chloride.

Part C: The crude 1-(chlorophenyl)-5-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carbonyl chloride is dissolved in tetrahydrofuran (THF) (10 mL). 2,3-Dihydro-1H-inden-2-ylamine (0.40 g, 3.00 mmol) is added and the resulting solution is stirred for 42 hours at 25 °C. The mixture is concentrated in vacuo and the residue is purified by preparative liquid chromatography to give pure 1-(4-chlorophenyl)-5-(2,4-dichlorophenyl)-N-(2,3-dihydro-1H-inden-2-yl)-1H-1,2,4-triazole-3-carboxamide (393 mg, 81 % yield). MS (ESI⁺) 485.6. ¹H-NMR (400 MHz, DMSO-d₆): 3.06 (dd, J = 16 and 8 Hz, 2H), 3.21 (dd, J = 16 and 8 Hz, 2H), 4.71-4.82 (m, 1H), 7.12-7.16 (m, 2H), 7.19-7.24 (m, 2H), 7.39 (br d, J ~ 8 Hz, 2H), 7.52 (br d, J ~ 8 Hz, 2H), 7.60 (dd, J = 8 and 2 Hz, 1H), 7.71 (d, J = 2 Hz, 1H), 7.79 (d, J = 8 Hz, 1H), 8.93-8.97 (m, 1H, NH).

Analogously were prepared the examples 20-43:

20. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(1-ethynylcyclohexyl)-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 473.3.
21. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(2-methylcyclohexyl)-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 465.5.
22. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(4-methylcyclohexyl)-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 465.5.
23. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-cyclooctyl-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 477.3.
24. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(azepan-1-yl)-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 466.4.
25. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-cycloheptyl-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 465.5.
26. N-t-Butyl-1-(4-chlorophenyl)-5-(2,4-dichlorophenyl)-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 425.4.
27. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(1,1-diethylprop-2-yn-1-yl)-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 461.5.
28. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(2,2,2-trifluoroethyl)-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 451.3.
29. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(exo-bicyclo[2.2.1]hept-2-yl)-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 461.5.
30. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(4-(2-propyl)piperazin-1-yl)-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 480.3. ¹H-NMR (400 MHz, DMSO-d₆): 1.00 (d, J = 7 Hz, 6H), 2.46-2.56 (m, 4H), 2.72 (septet, J = 7 Hz, 1H), 3.66-3.74 (m, 4H), 7.36 (br d, J = 8 Hz, 2H), 7.51 (br d, J = 8 Hz, 2H), 7.59 (dd, J = 8 and 2 Hz, 1H), 7.72 (d, J = 2 Hz, 1H), 7.75 (d, J = 8 Hz, 1H).

31. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(hexahydrocyclopenta[c]pyrrol-
2(1H)-yl)-1H-1,2,4-triazole-3-carboxamide. MS (ESI⁺) 476.4.
32. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-pentyl-1H-1,2,4-triazole-3-
carboxamide. MS (ESI⁺) 435.5.
- 5 33. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(2,2-dimethylpropyl)-1H-1,2,4-
triazole-3-carboxamide. MS (ESI⁺) 439.6.
34. 1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-N-(3-(trifluoromethyl)phenyl)-1H-
1,2,4-triazole-3-carboxamide. MS (ESI⁺) 511.7.
- 10 35. 1'-[1-(4-Chlorophenyl)-5-(2,4-dichlorophenyl)-1H-1,2,4-triazol-3-yl]carbonyl]-
1,4'-bipiperidine. MS (ESI⁺) 520.5.
36. 1-(4-Chlorophenyl)-N-(4-chlorophenyl)-5-(2,5-dichlorophenyl)-N-methyl-1H-
1,2,4-triazole-3-carboxamide. MS (ESI⁺) 491.4.
37. 1-(4-Chlorophenyl)-5-(2,5-dichlorophenyl)-N-(1-ethynylcyclohexyl)-1H-1,2,4-
triazole-3-carboxamide. MS (ESI⁺) 473.4.
- 15 38. 1-(4-Chlorophenyl)-5-(2,5-dichlorophenyl)-N-(2-methylcyclohexyl)-1H-1,2,4-
triazole-3-carboxamide. MS (ESI⁺) 465.5.
39. 1-(4-Chlorophenyl)-5-(2,5-dichlorophenyl)-N-(4-methylcyclohexyl)-1H-1,2,4-
triazole-3-carboxamide. MS (ESI⁺) 465.6.
40. 1-(4-Chlorophenyl)-5-(2,5-dichlorophenyl)-N-cyclooctyl-1H-1,2,4-triazole-3-
20 carboxamide. MS (ESI⁺) 477.3.
41. 1-(4-Chlorophenyl)-5-(2,5-dichlorophenyl)-N-cycloheptyl-1H-1,2,4-triazole-3-
carboxamide. MS (ESI⁺) 465.6.
42. 1-(4-Chlorophenyl)-5-(2,5-dichlorophenyl)-N-cyclopentyl-1H-1,2,4-triazole-3-
carboxamide. MS (ESI⁺) 435.5.
- 25 43. 1-(4-Chlorophenyl)-5-(2,5-dichlorophenyl)-N-(2,2-dimethylpropyl)-1H-1,2,4-
triazole-3-carboxamide. MS (ESI⁺) 439.6.

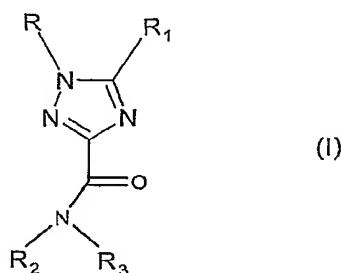
Pharmacological test results of a subset of the compounds of the invention, obtained with the assays described above, are given in the table below:

30

Example	Human cannabinoid-CB ₁ receptors	
	<i>In vitro</i> affinity <i>pK_i</i> value	<i>In vitro</i> antagonism <i>pA₂</i> value
Example 2	6.6	7.2
Example 3	6.9	8.7
Example 5	6.9	
Example 9	7.4	8.2
Example 11	6.3	

Claims

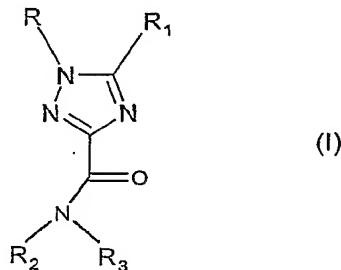
1. A method of treating disorders involving CB₁ cannabinoid neurotransmission such as psychosis, anxiety, depression, attention deficits, memory disorders, cognitive disorders, appetite disorders, obesity, addiction, appetite, drug dependence, neurodegenerative disorders, dementia, dystonia, muscle spasticity, tremor, epilepsy, multiple sclerosis, traumatic brain injury, stroke, Parkinson's disease, Alzheimer's disease, epilepsy, Huntington's disease, Tourette's syndrome, cerebral ischaemia, cerebral apoplexy, craniocerebral trauma, stroke, spinal cord injury, neuroinflammatory disorders, plaque sclerosis, viral encephalitis, demyelination related disorders, as well as for the treatment of pain disorders, including neuropathic pain disorders, septic shock, glaucoma, diabetes, cancer, emesis, nausea, gastrointestinal disorders, gastric ulcers, diarrhoea and cardiovascular disorders, characterized in that for the preparation of a pharmaceutical composition for the treatment of said disorders a compound of formula (I) is used



- 20 wherein
- R and R₁ independently represent a phenyl, naphtyl, thienyl, pyridyl, pyrimidyl, pyrazinyl, pyridazinyl or triazinyl group, which groups may be substituted with 1-4 substituents X, which can be the same or different, from the group branched or unbranched (C₁₋₃)-alkyl or alkoxy, hydroxy, halogen, trifluoromethyl, trifluoromethylthio, trifluoromethoxy, nitro, amino, mono- or dialkyl (C₁₋₂)-amino, mono- or dialkyl (C₁₋₂)-amido, (C₁₋₃)-alkoxycarbonyl, trifluoromethylsulfonyl, sulfamoyl, (C₁₋₃)-alkylsulfonyl, carboxyl, cyano, carbamoyl, (C₁₋₃)-dialkylaminosulfonyl, (C₁₋₃)-monoalkylamino-sulfonyl and acetyl,
- 25
- R₂ represents a hydrogen atom or a branched or unbranched C₁₋₈ alkyl or C₁₋₈ cycloalkyl-alkyl group or a phenyl, benzyl or phenethyl group which aromatic rings may be substituted with 1-4 substituents X, wherein X has the meaning as indicated above, or R₂ represents a pyridyl or thienyl group,
- 30
- R₃ represents branched or unbranched C₁₋₈ alkyl, C₁₋₈ alkoxy, C₃₋₈ cyclo-alkyl, C₅₋₁₀ bicycloalkyl, C₆₋₁₀ tricycloalkyl, C₃₋₈ alkenyl, C₅₋₈ cycloalkenyl, which
- 35

groups may optionally contain one or more heteroatoms from the group (O, N, S), which groups may be substituted with a hydroxy group, an ethynyl group or 1-3 fluoro atoms, or R₃ represents a phenyl, benzyl or phenethyl group which aromatic rings may be substituted with 1-4 substituents X, wherein X has the meaning as indicated above, or R₃ represents a pyridyl, pyrimidyl, pyrazinyl, pyridazinyl, triazinyl or thienyl group which heteroaromatic rings may be substituted with 1-2 substituents X, wherein X has the meaning as indicated above, or R₃ represents a group NR₄R₅ wherein R₄ and R₅, together with the nitrogen atom to which they are bonded, form a saturated or unsaturated, monocyclic or bicyclic, heterocyclic moiety having 4 to 10 ring atoms, which heterocyclic group contains one or two heteroatoms from the group N, O or S, which heteroatoms can be the same or different, which heterocyclic moiety may be substituted with a branched or unbranched C₁₋₃ alkyl, hydroxy or trifluoromethyl group or a fluoro atom, or R₂ and R₃, together with the nitrogen atom to which they are bonded, form a saturated or unsaturated, monocyclic or bicyclic, heterocyclic moiety having 4 to 10 ring atoms, which heterocyclic group contains one or two heteroatoms from the group N, O or S, which heteroatoms can be the same or different, which heterocyclic moiety may be substituted with a branched or unbranched C₁₋₃ alkyl, hydroxy, piperidinyl or trifluoromethyl group or a fluoro atom, and prodrugs, stereoisomers and salts thereof.

2. Compounds of the general formula (I)



25

wherein

- R and R₁ have the meanings as given in claim 1,
- R₂ represents a hydrogen atom or a branched or unbranched C₁₋₈ alkyl group
- R₃ represents branched or unbranched C₂₋₈ alkoxy, C₃₋₈ cycloalkyl, C₅₋₁₀ bicycloalkyl, C₆₋₁₀ tricycloalkyl, C₄₋₈ alkenyl, C₅₋₈ cycloalkenyl, which groups

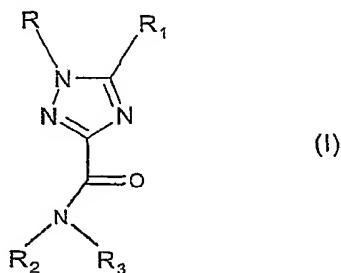
may optionally contain one or more heteroatoms from the group (O, N, S), which groups may optionally be substituted with a hydroxy group or 1-3 fluoro atoms, or R₃ represents a C₃₋₈ trifluoroalkyl or C₂₋₈ fluoroalkyl group, or R₃ represents a benzyl or phenethyl group which aromatic rings may be substituted with 1-4 substituents X, wherein X has the meaning as given in claim 1, or R₃ represents a 3-pyridyl, 4-pyridyl, pyrimidyl, pyrazinyl, pyridazinyl, triazinyl or thienyl group which heteroaromatic rings may be substituted with 1-2 substituents X, wherein X has the meaning as given in claim 1, or R₃ represents a group NR₄R₅ wherein

R₄ and R₅ together with the nitrogen atom to which they are bonded form a saturated or unsaturated, monocyclic or bicyclic, heterocyclic moiety having 4 to 10 ring atoms, which heterocyclic group contains one or two heteroatoms from the group N, O or S, which heteroatoms can be the same or different, which heterocyclic moiety may be substituted with a branched or unbranched C₁₋₃ alkyl, hydroxy or trifluoromethyl group or a fluoro atom, or

- R₂ and R₃, together with the nitrogen atom to which they are bonded, form a saturated or unsaturated, monocyclic or bicyclic, heterocyclic moiety having 4 to 10 ring atoms, which heterocyclic group contains one or two heteroatoms from the group N, O or S, which heteroatoms can be the same or different, which heterocyclic moiety may be substituted with a branched or unbranched C₁₋₃ alkyl, hydroxy, piperidinyl or trifluoromethyl group or a fluoro atom, with the proviso that this heterocyclic moiety is not an unsubstituted piperidinyl or unsubstituted morpholinyl group or 2,2,6,6-tetraalkylpiperidinyl group, and prodrugs, stereoisomers and salts thereof.

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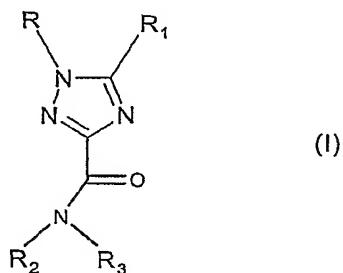
3. Compounds of the general formula (I)



wherein

- R and R₁ independently represent a phenyl, naphtyl, thienyl, pyridyl, pyrimidyl, pyrazinyl, pyridazinyl or triazinyl group, which groups are substituted with 1-4 substituents X, wherein X has the meaning as given in claim 1,
- R₂ and R₃ have the meanings as given in claim 2, and prodrugs, stereoisomers and salts thereof.

10 4. Compounds of the general formula (I)



wherein

- 15 R and R₁ independently represent a phenyl group which phenyl groups are substituted with 1-4 substituents which can be the same or different, from the group methyl, methoxy, halogen, trifluoromethyl or cyano or R and R₁ independently represent a phenyl, thienyl or pyridyl group, which phenyl group is substituted with 1-4 substituents, which can be the same or different, from the group methyl, methoxy, halogen, trifluoromethyl or cyano,
- 20 - R₂ has the meaning as given in claim 2,
- 25 - R₃ represents a group NR₄R₅ wherein R₄ and R₅ together with the nitrogen atom to which they are bonded form a saturated or unsaturated, monocyclic or bicyclic, heterocyclic moiety having 4 to 10 ring atoms, which heterocyclic group contains one or two heteroatoms from the group N, O or S, which heteroatoms can be the same or different, which heterocyclic moiety may be substituted with a branched or unbranched C₁₋₃ alkyl, hydroxy or trifluoromethyl group or a fluoro atom, and prodrugs, stereoisomers and salts thereof.
- 30

5. Pharmaceutical compositions containing a pharmacologically active amount of at least one compound of one of the claims 1-4 as an active ingredient.

6. Use of a compound of one of the claims 1-4 for the preparation of a pharmaceutical composition for the treatment of disorders involving cannabinoid neurotransmission.
5
7. Use as in claim 6 characterised in that said disorders are: psychosis, anxiety, depression, attention deficits, memory disorders, cognitive disorders, appetite disorders, obesity, addiction, appetence, drug dependence, neurodegenerative disorders, dementia, dystonia, muscle spasticity, tremor, epilepsy, multiple
10 sclerosis, traumatic brain injury, stroke, Parkinson's disease, Alzheimer's disease, epilepsy, Huntington's disease, Tourette's syndrome, cerebral ischaemia, cerebral apoplexy, craniocerebral trauma, stroke, spinal cord injury, neuroinflammatory disorders, plaque sclerosis, viral encephalitis, demyelinisation related disorders, as well as for the treatment of pain disorders, including
15 neuropathic pain disorders, septic shock, glaucoma, diabetes, cancer, emesis, nausea, gastrointestinal disorders, gastric ulcers, diarrhoea and cardiovascular disorders.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP 03/50628

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61K31/41 C07D249/10 A61P25/18

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A61K C07D A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, BEILSTEIN Data, CHEM ABS Data, BIOSIS, EMBASE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category ^a	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>CHEMICAL ABSTRACTS, vol. 85, no. 23, 6 December 1976 (1976-12-06) Columbus, Ohio, US; abstract no. 177322, HARHASH, ABDEL H. ET AL: "Reactions of 2-aryl-4-arylhydrazone-.DELTA.2-oxazolin-5 -ones with acid amides and diamines" XP002225874 RN's: 60776-63-8, 60776-64-9, 60776-65-0, 60776-66-1 abstract & INDIAN JOURNAL OF CHEMISTRY, SECTION B: ORGANIC CHEMISTRY INCLUDING MEDICINAL CHEMISTRY (1976), 14B(4), 268-72,</p> <p>---</p> <p>-/-</p>	2,3

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
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- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

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- *&* document member of the same patent family

Date of the actual completion of the international search

12 February 2004

Date of mailing of the international search report

01/03/2004

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl.
Fax: (+31-70) 340-3016

Authorized officer

Steendijk, M

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 03/50628

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	CHEMICAL ABSTRACTS, vol. 124, no. 17, 22 April 1996 (1996-04-22) Columbus, Ohio, US; abstract no. 232340, ELNAGDI, MOHAMED HILMY ET AL: "Studies with polyfunctionally substituted heteroaromatics: 2-phenyl-4-p-tolylhydrazone-2-oxazoline-5- one as a precursor for the synthesis of substituted 1,2,4-triazoles and pyridines" XP002225875 RN: 174682-65-6 abstract & HETEROATOM CHEMISTRY (1995), 6(6), 589-92, ---	2,3
Y	JAGEROVIC ET AL.: "A novle class of heterocyclic cannabinoids..." DRUGS OF THE FUTURE, vol. 27, no. suppl. A, September 2002 (2002-09), page 271 XP001121041 cited in the application page 271 ---	1-7
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Y	WO 01 58869 A (PANDIT CHENNAGIRI R ;SQUIBB BRISTOL MYERS CO (US); WROBLESKI STEPH) 16 August 2001 (2001-08-16) claim 3 ---	1-7
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Information on patent family members

International Application No

PCT/EP 03/50628

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 03/50628

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